

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

```
Run on:      June 29, 2002, 17:13:47 ; Search time 17398.4 Seconds
              (without alignments)
              1042.620 Million cell updates/sec
```

Title: US-09-303-518D-125
 Perfect score: 1344
 Sequence: 1 atgattaaatcaaaaaagg.....coattgaagaaggaggtga 1344

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 13736207 seqs, 6748477542 residues

Total number of hits satisfying chosen parameters: 27472414

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Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
                  Maximum Match 10%
                  Listing first 10%

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Database :

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Est:
1:  em_estba:*
2:  em_esthum:*
3:  em_estin:*
4:  em_estmu:*
5:  em_estov:*
6:  em_estpl:*
7:  em_estro:*
8:  em_htc:*
9:  gb_estli:*
10: gb_est2:*
11:  gb_htc:*
12:  gb_gss:*
13:  em_gss_hum:*
14:  em_gss_inv:*
15:  em_gss_pln:*
16:  em_gss_vrt:*

```

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query		Length	DB	ID	Description
		Match	%				
1	107	8.0		349	12	CNS07GYI	AL610380 Anopheles
2	39	2.9		684	10	BF728886	BF728886 1031102C0
3	38.8	2.9		686	9	AL628099	AL628099 AL628099
4	38.6	2.9		497	12	P947R	AL390645 Leishmani
5	38.2	2.8		455	10	B1956450	B1956450 HVSMER000
6	37.8	2.8		437	10	T44736	T44736 7999 Lambda
7	37.6	2.8		384	9	AL1392198	AL1392198 NCSMID6T3
8	37.6	2.8		528	10	BG606950	BG606950 WHE2492_E
9	37.6	2.8		540	10	BF260729	BF260729 HVSMER002
10	37.6	2.8		544	10	BF260723	BF260723 HVSMER002
11	37.6	2.8		551	10	BF253989	BF253989 HVSMER000
12	37.6	2.8		574	10	BF265938	BF265938 HV_CEA001
13	37.6	2.8		622	10	B1957371	B1957371 HVSMER000
14	37.6	2.8		634	10	B1959896	B1959896 HVSMER002
15	37.6	2.8		636	10	B1959473	B1959473 HVSMER001
16	37.6	2.8		697	10	BF628426	BF628426 HVSMER000
17	37.6	2.8		704	10	BF628832	BF628832 HVSMER000

RESULT	4
147R	
OCUS	
FINITION	
P947R	497 bp DNA linear
Leishmania major	Friedlin PAC P947 right end-sequence, genomic
	GSS 25-JUL-2000

JOURNAL
COMMENTJOURNAL
COMMENT

Clemson University
00 Jordan Hall, Clemson, SC 29634, USA

ORGANISM	<p>Arabidopsis thaliana Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons: core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.</p>
REFERENCE	<p>1 (bases 1 to 437)</p>
AUTHORS	<p>Newman,T., debruin,F.J., Green,P., Keegstra,K., Kende,H., McIntosh L., Ohlrogge,J., Raikhel,N., Somerville,S., Thomasnow,M., Retzel ,E. and Somerville,C.</p>
TITLE	<p>genes galore: a summary of methods for accessing results from large-scale partial sequencing of anonymous Arabidopsis cdna clones</p>
JOURNAL	<p>Plant Physiol. 106, 1241-1255, (1994)</p>
MEDLINE	<p>95148729</p>
COMMENT	<p>On Jan 7, 1998 this sequence version replaced gi:949002. Contact: Thomas Newman MSU-DOE Plant Research Laboratory Michigan State University MSU-DOE-PRL, Michigan State University, Plant Biology Bldg.,E. Lansing,Mi Tel: 517-353-0854 Fax: 517-353-9168 Email: 22313tcn@bm.cl.msu.edu See primer: T7.</p>

FEATURES	source	Location/Qualifiers
1.	.437	
/organism="Arabidopsis thaliana"		
/strain="var columbiana"		
/db_xref="taxon:3702"		
/clone="123P21n7"		
/clone_lib="Lambda-PRL2"		
/note="Vector: lambda Zip-Lox; Site_1: Sal; Site_2: Not; Lambda PRL2 is a cDNA library derived from equal quantities of 4 pools of mRNA. The mRNA sources were 1) 7 day germinated etiolated seedlings; 2) tissue culture grown roots; 3) staged plants half with 24 hour light cycle, half on 16 hr light, 8 hour dark- rosettes; 4) same plants as 3 but aerial tissue (stems, flowers and siliques). The vector is BRL's lambda zip_Lox. The cDNA inserts were directionally cloned with Sal-Not arms using oligo dt primed cDNA."		
99 a	87 c	103 g 132 t
BASE COUNT		16 others
ORIGIN		

	Query Match	2.8%;	Score 37.8;	DB 10;	Length 437;
	Best Local Similarity	55.8%;	Pred. No. 10;		
	Matches 72;	Conservative	Mismatches	57;	Indels 0; Gaps 0;
Qy	128	tgaagtcgaaggcgatgccctcaaaaaggccaagtgcgtgtttgaagacaaaaga	187		
Db	71	TCAAAGCTACACAGATCTCTGAATCGAAAAGCTTGAGCTGTGTTAATCGCGAGGAGA	130		
Qy	188	atccggggcgtggtttactgcgcgcgttcaggcaaaatccgcgcgtatccagctggcg	247		
Db	131	AATGGTGTGTATGTGTACGGCGCAGACTCCAACGAATATCGCGGTGATTAAATTTGGG	190		
Qy	248	aaaagcgcg	256		
Db	191	GAAGAAGAGAG	199		

RESULT	7
AI392198/c	
LOCUS	AI392198
DEFINITION	NCSMID6T3 Subtracted Mycelial Neurospora crassa cDNA clone SMID6 5 similar to dolichyl-phosphate-mannose synthase, mRNA sequence.
ACCESSION	AI392198
VERSION	AI392198.1 GI:4220005
KEYWORDS	EST.
SOURCE	Neurospora crassa.
ORGANISM	Neurospora crassa
REFERENCE	Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes; Sordariales; Sordariaceae; Neurospora. 1 (bases 1 to 384)

```

Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clmenson.edu
Total hq bases = 327
Seq primer: AATTACCCCTCACTAAAGGG
High quality sequence stop: 453.
Location/Qualifiers
1. .455
/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone="HVSME0003G23f"
/clone_lib="Hordeum vulgare rachis EST library HVCDNA0015
(normal)"
/tissue_type="rachis"
/lab_host="TJC121"
/notes="vector: pBluescript SK(-); Site_1: EcoRT; Site_2:
XhoI; Plants were grown at Washington State University,
Pullman, WA in a greenhouse, the rachises were excised and
frozen in liquid nitrogen (Kleinhofs lab). In the TJ Close
lab at the University of California, Riverside total RNA
was prepared, poly(A) was purified, one primary
unamplified cDNA library was made, and 1 million pfu were
in vivo excised to give pBluescript SK(-) cDNA phagemids
(Chn). Phagemids were plated and picked at the Clemson
University Genomics Institute (CUGI) (Begum, Palmer,
Frisch, Atkins and Wing). Plasmid DNA preparations, DNA
sequencing and sequence analysis were performed at CUGI
(Wing, Yu, Frisch, Henry, Simmons, Rambo, Main). The
sequence has been trimmed to remove vector sequence and
contains a minimum of 100 bases of phred value 20 or
above. For more details on library preparation and
sequence analysis see
http://www.genome.clemson.edu/projects/barley. To order
this clone see http://www.genome.clemson.edu/orders. Also
see Close TJ, Wing R, Kleinhofs A, Wise R (2001)
Genetically and physically anchored EST resources for
barley genomics, Barley Genetics Newsletter 31:29-30.
(http://wheat.pw.usda.gov/gpages/bgn/31/cover.html)."
83 a 150 c 150 g 2 others

```

	Query Match	2.8%;	Score 38.2;	DB 10;	Length 455;
	Best Local Similarity	49.5%;	Pred. No. 8.2;		
Matches	97; Conservative	0;	Mismatches	99;	Indels 0; Gaps
Qy	908	cgggtattgaacggcgcgattacacaaggcgcgcacgattatttgggacacatccacaatc	967		
Db	185	CGGTGATCTTCGACGCCGGGAAGACAGGGGCACGGCGCTGCACGCTCTTCGGGTTCGCACAAGA	244		
Qy	968	agatttcggtatcgataagaaggcgcgaagagcgtgttcgctcgtgggttgcgcgcgcgcgc	1027		
Db	245	AGATGGAGCTCGTCGACGCTGGCGGCACACATCGAGGCTCTTCGCCAAGGTGAGGCCGGGC	304		
Qy	1028	cggcaaatcctcatcacgcggtacaaacctcggccattcttcgtgaaaaacaaactcttca	1087		
Db	305	TGAGCTCGTAGCGGGACGGCCGANGAGGCCGCCAAGTCCATCACACCCCTCTGGAGA	364		
Qy	1088	agttcaacacagccgt	1103		

RESULT		6	T44736	437 bp	mRNA	linear	EST 07-JAN-1998
Locus			T44736				
DEFINITION			7999 Lambda-PRL2 Arabidopsis thaliana cDNA clone l23P2lT7, mRNA sequence.				
ACCESSION			T44736				
VERSION			T44736.1				
KEYWORDS			GI:2759537				
SOURCE			EST.				
			thale cress.				

BASE COUNT	ORIGIN	Genetically and physically anchored EST resources for barley genomics. Barley Genetics Newsletter 31:29-30. (http://wheat.pw.usda.gov/ggpages/bgn/31/cover.html)
110 a	165 c	174 g 92 t 3 others.

BASE COUNT	110 a	165 c	174 g	92 t	3 others.	
ORIGIN	{http://wheat.pw.usda.gov/ggpages/bgn/31/cover.html}	"				
Query Match	2.8%	Score 37.6;	DB 10;	Length 544;		
Best Local Similarity	49.5%	Pred. No. 13;				
Matches	97;	Conservative	0;	Mismatches	99;	Indels
					0;	Gaps
Qy	908	cggatttgacgcgcgattacacaaaggcgcgcacgattatttgggacgtctaccacaatc	967			
Db	167	CGGTGATCTTCACGCCCGGAAGACACGGGCGACGCGCGTGCACGCTTCCTCGGTTTCGACAA	226			
Qy	968	agatttcggtatcgaaagaaggcgcgcgacaaagactgttcgctgggtgcgcgcagc	1027			
Db	227	AGATGGAGCTCTGCAGCTCGGCCACCAATCAGAGTCTTCGCCAAAGGTGGAGCGGGCG	286			
Qy	1028	cggacaataactccatcacgcgtacacccctcgccatttctctgaaaaaacaacttcca	1087			
Db	287	TGAGTCTGTACGCCGGACGGCCGACGAGGCCCGCAAGTCCATCACACCCCTGCTGGAGA	346			
Qy	1088	agttcaacacagccgt	1103			
Db	347	AGGCCAAGAGCGCGT	362			

[illegible]

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FEATURES
source
High quality sequence stop: 540.
Location/Qualifiers
1. .551
/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone_lib="Hordeum vulgare seedling root EST library
HVCN0A0007 (Etiolated and unstressed)"
/tissue_type="Seedling root"
/lab_host="TJC121"
/notes="Vector: lambdaZAP; Site_1: EcoRI; Site_2: XhoI;
Seeds were surface sterilized then germinated under axenic
conditions in the dark at room temperature on filter paper

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Query Match	2.8%	Score 37.6;	DB 10;	Length 551;
Best Local Similarity	49.5%	Pred. No.13;		
Matches 97;	Conservative 0;	Mismatches 99;	Indels 0;	Gaps 0;
QY 908	cgggtattgaacg	cgcgattacacaa	ggcgcgacgattatt	ggagcgtaccacaatc 967
Db 25	CGGTGATCTTC	AGCCCGGAGAC	ACGGGCGACGGCGGTGCACGCTT	CCGGTTCGACAAGA 84
QY 968	agatttcggtatc	gaagaagcgcgac	gagcaaaagctgttc	gcgtggttcgcccgcagc 1027
Db 85	AGATGAGCTGT	CGACGTCGGCG	ACGACATCGAGGTCTCC	CAAGGTGAGCCGGGGC 144
QY 1028	cggacaataatc	atcacgcgtacac	accctcgccatttc	cttgtaaaacaaactttca 1087
Db 145	TCAGTCTGTAG	CGCGGCGCGGAG	GCGCCCAAGTCCATC	ACACCCCTGCTGGAGA 204
QY 1088	agttcaacacag	cgcgt 1103		
Db 205	AGGCCAAGAG	CGCGGT 220		

RESULT	12	
LOCUS	BF265938	
DEFINITION	HV_CEA0013L17f Hordeum vulgare seedling green leaf EST library	
	574 bp	linear EST 23-OCT-2001
	HVCDA0004	(Blumeria challenged)
	HV_CEA0013L17f	Hordeum vulgare cDNA clone
ACCESSION	BF265938	
VERSION	BF265938.2	GI:13262431
KEYWORDS	EST.	
SOURCE	barley.	
ORGANISM	Hordeum vulgare	
	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;	
	Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae;	
	1 (bases 1 to 574)	
REFERENCE	Wing, R., Close, T.J., Kleinhofs, A., Wise, R., Wei, F., Begum, D., Frisch, D., Yu, Y., Henry, D., Palmer, M., Rambo, T., Simmons, J., Choi, D.W., Fenton, R.D., Oates, R. and Main, D.	
AUTHORS	Development of a genetically and physically anchored EST resource for barley genomics: Blumeria infected incompatible (Mial3) seedling leaf cDNA library	
TITLE	Unpublished (2001)	
JOURNAL	On Nov 17, 2000 this sequence version replaced gi:11196932.	
COMMENT	Contact: Wing RA Clemson University Clemson University	


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|||||
Db 245 AGATGGAGCTCGTGGAGCGGACGACATCGAGGTCTTCCCAAGGTGGAGCGGGGC 304
QY 1028 cggacaaatactcaccacgcgtacacacccctcgccattctctgaaacaaactcttca 1087
Db 305 TGAGCTGTGATCGGCGGACGCGGAGGAGGCGGCAAGTCCATCATCACCCCTGCTGGAGA 364
QY 1088 agttcaacacagccgt 1103
Db 365 AGGCCAAGAGCGCGGT 380

RESULT 14
BI959896
LOCUS
DEFINITION
HVSME0022E12f Hordeum vulgare rachis EST library HVCDNA0015
(normal) Hordeum vulgare cDNA clone HVSME0022E12f, mRNA sequence.
ACCESSION
BI959896
VERSION
BI959896.1 GI:16311151
KEYWORDS
EST.
SOURCE
barley.
ORGANISM
Hordeum vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
; Triticeae; Hordeum.
1 (bases 1 to 634)
Wing, R., Close, T.J., Kleinhofs, A., Wise, R., Chin, A., Begum, D.,
Frisch, D., Atkins, M., Yu, Y., Henry, D., Palmer, M., Rambo, T., Simmons
J., Oates, R. and Main, D.
Development of a genetically and physically anchored EST resource
for barley genomics: Morex rachis cDNA library
Unpublished (2001)
Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Total hq bases = 465
Seq primer: AATTAACCCCTCACTAAAGGG
High quality sequence stop: 605.
Location/Qualifiers
1. 634
/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone="HVSME0022E12f"
/clone_lib="Hordeum vulgare rachis EST library HVCDNA0015
(normal)"
/tissue_type="Rachis"
/lab_host="TJCI21"
/notes="Vector: pBluescript SK(-); Site 1: EcoRI; Site 2:
XhoI; Plants were grown at Washington State University,
Pullman, WA in a greenhouse, the rachises were excised and
frozen in liquid nitrogen (Kleinhofs lab). In the TJ Close
lab at the University of California, Riverside total RNA
was prepared, poly(A) was purified, one primary
unamplified cDNA library was made, and 1 million pfu were
in vivo excised to give pBluescript SK(-) cDNA phagemids
(Chin). Phagemids were plated and picked at the Clemson
University Genomics Institute (CUGI) (Begum, Palmer,
Frisch, Atkins and Wing). Plasmid DNA preparations, DNA
sequencing and sequence analysis were performed at CUGI
(Wing, Yu, Frisch, Henry, Simmons, Rambo, Main). The
sequence has been trimmed to remove vector sequence and
contains a minimum of 100 bases of phred value 20 or
above. For more details on library preparation and
sequence analysis see
http://www.genome.clemson.edu/projects/barley. To order
this clone see http://www.genome.clemson.edu/orders
see Close TJ, Wing R, Kleinhofs A, Wise R (2001)
Genetically and physically anchored EST resources for

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barley genomics. Barley Genetics Newsletter 31:29-30.
(http://wheat.pw.usda.gov/ggpages/bgn/31/cover.html)."
BASE COUNT 131 a 185 c 205 g 113 t
ORIGIN

Query Match 2.8%; Score 37.6; DB 10; Length 634;
Best Local Similarity 49.5%; Pred. No. 14;
Matches 97; Conservative 0; Mismatches 99; Indels 0; Gaps 0;

QY 908 cggattgaacgagcgattacacaaaggcgagcattattgggagcgtaccacatc 967
|||||
Db 177 CGGTGATCTTCAGCCGCGGAAACACGCGGCGGCGGTTCGGTTCGACAAGA 236
QY 968 agatttccttattcgaagaagccgcagcagacttctcggttggtcgccgcagc 1027
|||||
Db 237 AGATGGAGCTCGTCCAGCTCGCGGACGACATCGAGGTCTTCCCAAGTGGAGCGGGGC 296
QY 1028 cggacaaatactcaccacgcgtacacacccctcgccattctctgaaacaaactcttca 1087
|||||
Db 297 TGAGCTCGTACCGCGGACGCGGAGGAGGCGGCAAGTCCATCATCACCCCTGCTGGAGA 356
QY 1088 agttcaacacagccgt 1103
|||||
Db 357 AGGCCAAGAGCGCGGT 372

RESULT 15
BI959473
LOCUS
DEFINITION
HVSME0019N03f Hordeum vulgare rachis EST library HVCDNA0015
(normal) Hordeum vulgare cDNA clone HVSME0019N03f, mRNA sequence.
ACCESSION
BI959473
VERSION
BI959473.1 GI:16310728
KEYWORDS
EST.
SOURCE
barley.
ORGANISM
Hordeum vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
; Triticeae; Hordeum.
1 (bases 1 to 636)
Wing, R., Close, T.J., Kleinhofs, A., Wise, R., Chin, A., Begum, D.,
Frisch, D., Atkins, M., Yu, Y., Henry, D., Palmer, M., Rambo, T., Simmons
J., Oates, R. and Main, D.
Development of a genetically and physically anchored EST resource
for barley genomics: Morex rachis cDNA library
Unpublished (2001)
Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Total hq bases = 585
Seq primer: AATTAACCCCTCACTAAAGGG
High quality sequence start: 5
High quality sequence stop: 624.
Location/Qualifiers
1. 636
/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone="HVSME0019N03f"
/clone_lib="Hordeum vulgare rachis EST library HVCDNA0015
(normal)"
/tissue_type="Rachis"
/lab_host="TJCI21"
/notes="Vector: pBluescript SK(-); Site 1: EcoRI; Site 2:
XhoI; Plants were grown at Washington State University,
Pullman, WA in a greenhouse, the rachises were excised and
frozen in liquid nitrogen (Kleinhofs lab). In the TJ Close
lab at the University of California, Riverside total RNA

```

total hq bases = 340
Seq primer: AATTAAACCTCTACTTAAGGG
High quality sequence stop: 681.
Location/Qualifiers
1. 697
/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone="HVSMEB0006A02f"
/clone_lib="Hordeum vulgare seedling shoot EST library
HVCDNA0002 (Dehydration stress)"
/tissue_type="Seedling shoot"
/lab_host="TJC121"
/note="Vector: lambdaZAP; Site1: EcoRI; Site2: XhoI;
Seeds were surface sterilized then germinated under axenic
conditions in the dark at room temperature on filter paper
with water, nystatin and cefotaxime in covered
crystallization dishes. Five-day old seedlings were
incubated at 90% RH for 24 hr. Shoots were then harvested,
total RNA was prepared, poly(A) RNA was purified, one
primary unamplified cDNA library was made, 600000 pfu were
in vivo excised to give plusscript SK(-) cDNA phagemids.
These steps were performed in the TJ Close laboratory at
the University of California, Riverside (Choi, Close,
Penton). Phagemids were plated and picked at the Clemson
University Genomics Institute (CUGI) (Begum, Palmer,
Frisch, Atkins and Wing). Plasmid DNA preparations, DNA
sequencing and sequence analysis were performed at CUGI
(Wing, Yu, Frisch, Henry, Simmons, Oates, Rambo, Main).
The sequence has been trimmed to remove vector sequence
and contains a minimum of 100 bases of phred value 20 or
above. For more details on library preparation and
sequence analysis see
<http://www.genome.clemson.edu/projects/barley>. To order
this clone see <http://www.genome.clemson.edu/orders> Also
see Close TJ, Wing K, Kleinhoofs A, Wise R (2001)
Genetically and physically anchored EST resources for
barley genomics. Barley Genetics Newsletter 31:29-30.
(<http://wheat.pw.usda.gov/ggpages/bgn/31/cover.html>)"
206 c 126 t 1 others

Query Match	2.88;	Score 37.6;	DB 10;	Length 697;
Best Local Similarity	49.5%;	Pred No. 15;		
Matches 97;	Conservative	0;	Mismatches	99;
			Indels	0;
			Gaps	0;
QY	908	cggattgaacggcggattacacaaaggcgcgcagattatttggagcgtcaccaatc	967	
Db	169	CGGTGATCTTCAGCGCGGAGACACGCGCAGCGCGTGCAGCTTCCGTTTCACAGA	228	
QY	968	agatttcggttatcgaaagaagcgcgcagcagaagctgttcggctgggtgcgcgcagc	1027	
Db	229	AGATGAGCTGCTGCACGTCCGCGACGACATPCGAGGCTTCGCGCAAGTGGAGCCGGGC	288	
QY	1028	cggacaaatactccatcacgcgtacaaacccctgcgcattctctgaaaaaacaactctca	1087	
Db	289	TGAGTCTGTACGCCGCGCGGAGGAGCGCGCAAGTCATCACACCCTCTCTGGAGA	348	
QY	1088	gattceaceacgcgt	1103	
Db	349	AGGCTAAAGCCGCT	364	

RESULT	17	
BF628632		
LOCUS	704 bp	linear
DEFINITION	HVSMEB0006N12f Hordeum vulgare seedling shoot EST library	
	HVSMEB00002 (dehydration stress) Hordeum vulgare cDNA clone	
	HVSMEB0006N12f, mRNA sequence.	
ACCESSION	BF628632	
VERSION	BF628632.2	GI:13090280
KEYWORDS	EST.	
		EST 22-OCT-2000

JOURNAL
COMMENT

Unpublished (2001)
On Dec 19, 2000 this sequence version replaced gi:11892584.
Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwinc@clemson.edu

SOURCE
ORGANISM

barley.
Hordeum vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
; Triticeae; Hordeum.

REFERENCE
AUTHORS

Wing, R., Close, T.J., Kleinhofs, A., Wise, R., Begum, D., Frisch, D., Yu
, Y., Henry, D., Palmer, M., Rambo, T., Simmons, J., Choi, D.W., Fenton
, R.D., Oates, R. and Main, D.
Development of a genetically and physically anchored EST resource
for barley genomics: Morex drought-stressed seedling shoot cDNA
library

TITLE

Unpublished (2001)
On Dec 19, 2000 this sequence version replaced gi:11892790.

JOURNAL
COMMENT

Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Total hq bases = 580
Seq primer: AATTACCCCTCACTAAAGGG
High quality sequence stop: 663.

FEATURES
Source

Location/Qualifiers
1..704
/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone="HVSMEB0006N12f"
/clone_lib="Hordeum vulgare seedling shoot EST library
HVCNDA0002 (Dehydration stress)"
/tissue_type="Seedling shoot"
/lab_host="TJ121"
/notes="Vector: lambdaZAP; Site.1: EcoRI; Site.2: XhoI;
Seeds were surface sterilized then germinated under axenic
conditions in the dark at room temperature on filter paper
with water, nystatin and ceftaxime in covered
crystallization dishes. Five-day old seedlings were
incubated at 90% RH for 24 hr. Shoots were then harvested,
total RNA was prepared, poly(A) RNA was purified, one
primary unamplified cDNA library was made, 600000 pfu were
in vivo excised to give pBluescript SK(-) cDNA phagemids.
These steps were performed in the TJ Close laboratory at
the University of California, Riverside (Choi, Close,
Fenton). Phagemids were plated and picked at the Clemson
University Genomics Institute (CUGI) (Begum, Palmer,
Frisch, Atkins and Wing). Plasmid DNA preparations, DNA
sequencing and sequence analysis were performed at CUGI
(Wing, Yu, Frisch, Henry, Simmons, Oates, Rambo, Main).
The sequence has been trimmed to remove vector sequence
and contains a minimum of 100 bases of phred value 20 or
above. For more details on library preparation and
sequence analysis see
http://www.genome.clemson.edu/projects/barley. To order
this clone see http://www.genome.clemson.edu/orders Also
see close TJ, Wing R, Kleinhofs A, Wise R (2001)
Genetically and physically anchored EST resources for
barley genomics. Barley Genetics Newsletter 31:29-30.
(http://wheat.pw.usda.gov/gppages/bgn/31/cover.html)"

BASE COUNT
ORIGIN

146 a 206 c 226 g 126 t

Query Match 2.8%; Score 37.6; DB 10; Length 704;
Best Local Similarity 49.5%; Pred. No. 15;
Matches 97; Conservative 0; Mismatches 99; Indels 0; Gaps 0;

QY 908 cggattgaacgagcgattacacagcgccgcacgattatttggagcgtacacacac 967

Db 164 CGGTGATCTTCGACGCGGACACGCGCGGTCCAGCTCTTCGGTTCGACAAGA 223

QY 968 agatttcgattcagaagaagcgccgacgaagagctgttcgctgggtgctgcgcgcg 1027

Db 224 AGATGAGCTCTCTCAGCTCGGCGAGACATCGAGGTCTTCGCCAAGGTGGAGCGGGC 283
QY 1028 cggacaataactccatcacggtgtaacacccctcgccatttctgtaaaaaacaaactctca 1087
Db 284 TGAGTCTGTACGCGGAGCGCGCGCAGGAGCGCCAGTCCATCACACCCCTGCTGGAGA 343
QY 1088 agttcaacacagccat 1103
Db 344 AGCCCAAGAGCGCCCT 359

RESULT 18
BF259476

LOCUS
DEFINITION
759 bp mRNA linear EST 22-OCT-2001
HVSMEF0019D11f Hordeum vulgare seedling root EST library HVCNDA0007
(Etiolated and unstressed) Hordeum vulgare cDNA clone
HVSMEF0019D11f, mRNA sequence.

ACCESSION

BF259476

VERSION

BF259476.2 GI:131119939

KEYWORDS

EST.

SOURCE

barley.

ORGANISM

Hordeum vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
; Triticeae; Hordeum.

REFERENCE
AUTHORS

Wing, R., Close, T.J., Kleinhofs, A., Wise, R., Begum, D., Frisch, D., Yu
, Y., Henry, D., Palmer, M., Rambo, T., Simmons, J., Choi, D.W., Fenton
, R.D., Oates, R. and Main, D.
Development of a genetically and physically anchored EST resource
for barley genomics: Morex unstressed seedling root cDNA library
Unpublished (2001)

TITLE

On Nov 16, 2000 this sequence version replaced gi:11188505.
Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Total hq bases = 469
Seq primer: AATTACCCCTCACTAAAGGG
High quality sequence stop: 663.

JOURNAL
COMMENT

Location/Qualifiers
1..759
/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone="HVSMEF0019D11f"
/clone_lib="Hordeum vulgare seedling root EST library
HVCNDA0007 (Etiolated and unstressed)"
/tissue_type="Seedling root"
/lab_host="TJ121"
/notes="Vector: lambdaZAP; Site.1: EcoRI; Site.2: XhoI;
Seeds were surface sterilized then germinated under axenic
conditions in the dark at room temperature on filter paper
with water, nystatin and ceftaxime in covered
crystallization dishes. Five-day old seedling roots were
then harvested, total RNA was prepared, poly(A) RNA was
purified, one primary unamplified cDNA library was made,
and 1 million pfu were in vivo excised to give pBluescript
SK(-) cDNA phagemids. These steps were performed in the TJ
Close laboratory at the University of California,
Riverside (Choi, Close, Fenton). Phagemids were plated and
picked at the Clemson University Genomics Institute (CUGI)
(Begum, Palmer, Frisch, Atkins and Wing). Plasmid DNA
preparations, DNA sequencing and sequence analysis were
performed at CUGI (Wing, Yu, Frisch, Henry, Simmons, Oates
, Rambo, Main). The sequence has been trimmed to remove
vector sequence and contains a minimum of 100 bases of
phred value 20 or above. For more details on library
preparation and sequence analysis see

FEATURES
Source

Location/Qualifiers
1..759
/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone="HVSMEF0019D11f"
/clone_lib="Hordeum vulgare seedling root EST library
HVCNDA0007 (Etiolated and unstressed)"
/tissue_type="Seedling root"
/lab_host="TJ121"
/notes="Vector: lambdaZAP; Site.1: EcoRI; Site.2: XhoI;
Seeds were surface sterilized then germinated under axenic
conditions in the dark at room temperature on filter paper
with water, nystatin and ceftaxime in covered
crystallization dishes. Five-day old seedling roots were
then harvested, total RNA was prepared, poly(A) RNA was
purified, one primary unamplified cDNA library was made,
and 1 million pfu were in vivo excised to give pBluescript
SK(-) cDNA phagemids. These steps were performed in the TJ
Close laboratory at the University of California,
Riverside (Choi, Close, Fenton). Phagemids were plated and
picked at the Clemson University Genomics Institute (CUGI)
(Begum, Palmer, Frisch, Atkins and Wing). Plasmid DNA
preparations, DNA sequencing and sequence analysis were
performed at CUGI (Wing, Yu, Frisch, Henry, Simmons, Oates
, Rambo, Main). The sequence has been trimmed to remove
vector sequence and contains a minimum of 100 bases of
phred value 20 or above. For more details on library
preparation and sequence analysis see

Contact: Wing RA
Clemson University Genomics Institute
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Total hq bases = 476
Seq primer: AATTAACCCCTCACTAAAGGG
High quality sequence stop: 698.
Location/Qualifiers
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FEATURES
source

/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone_lib="HVSMB00001D17f"
HVCNDA0002 (Hordeum vulgare seedling shoot EST library
/tissue_type="Seedling shoot"
/lab_host="TJC121"
/note="Vector: lambdaZAP; Site_1: EcoRI; Site_2: XhoI;
Seeds were surface sterilized then germinated under axenic
conditions in the dark at room temperature on filter paper
with water, nystatin and cefotaxime in covered
crystallization dishes. Five-day old seedlings were
incubated at 90% RH for 24 hr. Shoots were then harvested,
total RNA was prepared, poly(A) RNA was purified, one
primary unamplified cDNA library was made, 600000 pfu were
in vivo excised to give pBluescript SK(-) cDNA phagemids.
These steps were performed in the TJ Close laboratory at
the University of California, Riverside (Choi, Close,
Fenton). Phagemids were plated and picked at the Clemson
University Genomics Institute (CUGI) (Begum, Palmer,
Frisch, Atkins and Wing). Plasmid DNA preparations, DNA
sequencing and sequence analysis were performed at CUGI
(Wing, Yu, Frisch, Henry, Simmons, Oates, Rambo, Main).
The sequence has been trimmed to remove vector sequence
and contains a minimum of 100 bases of phred value 20 or
above. For more details on library preparation and
sequence analysis see
http://www.genome.clemson.edu/projects/barley. To order
this clone see http://www.genome.clemson.edu/orders/Also
see Close TJ, Wing R, Kleinohs A, Wise R (2001)
Genetically and physically anchored EST resources for
barley genomics. Barley Genetics Newsletter 31:29-30.
(http://wheat.pw.usda.gov/ggpages/bgn/31/cover.html)"

BASE COUNT 177 a 220 c 252 g 134 t

Query Match 2.8%; Score 37.6; DB 10; Length 783;
Best Local Similarity 49.5%; Pred. No. 16;
Matches 97; Conservative 0; Mismatches 99; Indels 0; Gaps 0;

QY 908 cggattgaacgagcgattacacagcgccgacgattattggagcgatcacacaatc 967
Db 178 CGGTGATCTTCGACCGGGAACACGCGCGCGTCCAGCTTCCGGTTCGACAAAG 237
QY 968 agatttcggtatcgaagaagccgacgagcgtgttgcgtgggtgagccgcgcg 1027
Db 238 AGATGAGGCTGCTGACGCTCGCGGACGACATCGAGGTCTTCGCCAAGTGGAGCCGGGC 297
QY 1028 cggacaaaactccatcacgcgtacacacctcggcattcttcgaaacaaactctta 1087
Db 298 TGAGTCTGTCAGCGGCGGCGGAGGAGGCGCCGCAAGTCCATCATCACCCCTGCTGGAGA 357
QY 1088 agttcaacacagcgt 1103
Db 358 AGGCCAAGAGCGCGT 373

RESULT 21
BF254527

LOCUS
DEFINITION
BF254527
HVSMEF0004E12f Hordeum vulgare seedling root EST library HVCNDA0007
(Etioolated and unstressed) Hordeum vulgare cDNA clone
HVSMEF0004E12f, mRNA sequence.
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Barley.
Hordeum vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
; Triticeae; Hordeum.
REFERENCE
AUTHORS
1 (bases 1 to 841)
Wing, R., Close, T.J., Kleinohs, A., Wise, R., Begum, D., Frisch, D., Yu
Y., Henry, D., Palmer, M., Rambo, T., Simmons, J., Choi, D.W., Fenton
R.D., Oates, R. and Main, D.
Development of a genetically and physically anchored EST resource
for barley genomics: Morex unstressed seedling root cDNA library
Unpublished (2001)
On Nov 16, 2000 this sequence version replaced gi:11183632.
COMMENT
Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Total hq bases = 416
Seq primer: AATTAACCCCTCACTAAAGGG
High quality sequence stop: 580.
Location/Qualifiers
1...841

FEATURES
source

/organism="Hordeum vulgare"
/cultivar="Morex"
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/clone_lib="HVSMEF0004E12f"
HVCNDA0007 (Etioolated and unstressed)
/tissue_type="Seedling root"
/lab_host="TJC121"
/note="Vector: lambdaZAP; Site_1: EcoRI; Site_2: XhoI;
Seeds were surface sterilized then germinated under axenic
conditions in the dark at room temperature on filter paper
with water, nystatin and cefotaxime in covered
crystallization dishes. Five-day old seedling roots were
then harvested, total RNA was prepared, poly(A) RNA was
purified, one primary unamplified cDNA library was made,
and 1 million pfu were in vivo excised to give pBluescript
SK(-) cDNA phagemids. These steps were performed in the TJ
Close laboratory at the University of California,
Riverside (Choi, Close, Fenton). Phagemids were plated and
picked at the Clemson University Genomics Institute (CUGI)
(Begum, Palmer, Frisch, Atkins and Wing). Plasmid DNA
preparations, DNA sequencing and sequence analysis were
performed at CUGI (Wing, Yu, Frisch, Henry, Simmons, Oates
, Rambo, Main). The sequence has been trimmed to remove
vector sequence and contains a minimum of 100 bases of
phred value 20 or above. For more details on library
preparation and sequence analysis see
http://www.genome.clemson.edu/projects/barley. To order
this clone see http://www.genome.clemson.edu/orders/Also
see Close TJ, Wing R, Kleinohs A, Wise R (2001)
Genetically and physically anchored EST resources for
barley genomics. Barley Genetics Newsletter 31:29-30.
(http://wheat.pw.usda.gov/ggpages/bgn/31/cover.html)"

BASE COUNT 192 a 219 c 298 g 132 t

Query Match 2.8%; Score 37.6; DB 10; Length 841;
Best Local Similarity 49.5%; Pred. No. 16;
Matches 97; Conservative 0; Mismatches 99; Indels 0; Gaps 0;

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930	AGATGGAGCTCGTCGACCTCGGCGACGACATCGAGGTCTTCGCCAAGGTGAGCGCGGGC	299
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947		
948	agatttcggttatctgaagaagcgcgcaagagcgtcttcggttggttgcgcgcagc	1027
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RESULT	22
B1957008	
LOCUS	linear
DEFINITION	847 bp mRNA EST 22-OCT-2001 HVSME0006006orf Hordeum vulgare rachis EST library HVCDNA0015 (normal) Hordeum vulgare cDNA clone HVSME0006006orf, mRNA sequence.
ACCESSION	B1957008
VERSION	GI:16308261
KEYWORDS	
SOURCE	barley.
ORGANISM	Hordeum vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae ; Triticeae; Hordeum. 1 (bases 1 to 847) Wing, R., Close, T.J., Kleinhofs, A., Wise, R., Chin, A., Begum, D., Frisch, D., Atkins, M., Yu, Y., Henry, D., Palmer, M., Rambo, T., Simmons, J., Oates, R. and Main, D. Development of a genetically and physically anchored EST resource for barley genomics: Morex rachis cDNA library	
REFERENCE	
AUTHORS	
TITLE	

Journal Comment

Unpublished (2001)
Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Total hq bases = 592
Seq primer: AATTAAACCTCACTAAAGGG
High quality sequence stop: 667.

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1. 847
Location/Qualifiers
/organism="Hordeum vulgare"
/cultivar="Morex"
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(normal)"
/tissue_type="Rachis"
/lab_host="TJC121"
/Note="Vector: pBluescript SK(-); Site.1: EcoRI; Site.2:
XhoI; Plants were grown at Washington State University,
Pullman, WA in a greenhouse, the rachises were excised and
frozen in liquid nitrogen (Kleinhofs lab). In the TJ Close
lab at the University of California, Riverside total RNA
was prepared, poly(A) was purified, one primary
unamplified cDNA library was made, and 1 million pfu were
in vivo excised to give pBluescript SK(-) cDNA phagemids
(Chin). Phagemids were plated and picked at the Clemson
University Genomics Institute (CUGI) (Begum, Palmer,
Frisch, Atkins and Wing). Plasmid DNA preparations, DNA
sequencing and sequence analysis were performed at CUGI
(Wing, Xu, Frisch, Henry, Simmons, Rambo, Main). The
sequence has been trimmed to remove vector sequence and
contains a minimum of 100 bases of phred value 20 or
above. For more details on library preparation and

```

sequence analysis see
http://www.genome.clemson.edu/projects/barley. To order
this clone see http://www.genome.clemson.edu/orders Also
see Close TJ, Wing R, Kleinhofs A, Wise R (2001)
Genetically and physically anchored EST resources for
barley genomics. Barley Genetics Newsletter 31:29-30.
(http://wheat.pw.usda.gov/ggpages/bgn/31/cover.html)*
183 a 249 c 258 a 157 t

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99	0.000	0.000
100	0.000	0.000

	Query Match	2.8%	Score 37.6;	DB 10;	Length 847;
	Best Local Similarity	49.5%;	Pred. No. 16;		
	Matches	97;	Conservative	0;	Mismatches
				99;	Indels
					0;
					Gaps
					0;
QY	908	cggattgaacggcgcgattacacaaggcgcgacgattatttgggacgctaccacaatc	967		
Db	180	CGGTGNTCTTCACGCGGGAACACACGGCACGCGGTGCACGCTCTTCGGTTCGACAAGA	239		
QY	968	agatttcogtattcaagaagaagcgacgaagaagacttcgctgggttcgcccgcgacg	1027		
Db	240	AGATGGAGTCTGCTGACGTCGCGACGACATCGAGGTCTTCGCCAAGTGGAGCCGGGCG	299		
QY	1028	cggacaaatattccattcacgcggtacaacctcggcgacttcctctgaaaaacaaacttcca	1087		
Db	300	TGAGTCTGTACGCGGACGCGCCGACAGGAGCGCCCAAGTCCATCATACCCCTGCTGGAGA	359		
QY	1088	agttcaacacagccgt	1103		
Db	360	AGGCCAAGAGCGCGT	375		

RESULT 23

BI957786	BI957786	864 bp	linear	EST 22-OCT-2001
LOCUS	HVSMEn0011F07f	Hordeum vulgare	rachis	EST library HVCDNA0015
DEFINITION	(normal)	Hordeum vulgare	cDNA clone	HVSMEn0011F07f, mRNA sequence.
ACCESSION	BI957786			
VERSION	BI957786.1	GI:16309053		
KEYWORDS	EST.			
SOURCE	barley.			

ORGANISM	REFERENCE
<i>Hordeum vulgare</i>	Wing, R., Close, T.J., Kleinhofs, A., Wise, R., Chin, A., Begum, D., Frisch, D., Atkins, M., Yu, Y., Henry, D., Palmer, M., Rambo, T., Simmons, J., Oates, R. and Main, D.
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooidaeae	Development of a genetically and physically anchored EST resource for barley genomics: Morex rachis cDNA library
; Triticeae; Hordeum.	Unpublished (2001)

COMMENT

Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: twing@clemson.edu
Total hq bases = 458
Seq primer: AATTACCCCTCACTAAAGG
High quality sequence stop: 641.

FEATURES

```

source
1. .864
/organism="Hordeum vulgare"
/cultivar="Morex"
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/clone="HVSME0011F07"
/clone_lib="Hordeum vulgare rachis EST library HVCDNA0015
(normal)"
/tissue_type="Rachis"
/lab_host="TJCl21"
/note="Vector: pBluescript SK(-); Site_1: EcoRI; Site_2:

```


XhoI; Plants were grown at Washington State University, Pullman, WA in a greenhouse, the rachises were excised and frozen in liquid nitrogen (Kleinhoofs lab). In the TX Close lab at the University of California, Riverside total RNA was prepared, poly(A) was purified, one primary unamplified cDNA library was made, and 1 million pfu were in vivo excised to give pBluescript SK(-) cDNA phagemids (Chin). Phagemids were plated and picked at the Clemson University Genomics Institute (CUGI) (Begum, Palmer, Frisch, Atkins and Wing). Plasmid DNA preparations, DNA sequencing and sequence analysis were performed at CUGI (Wing, Yu, Frisch, Henry, Simmons, Rambo, Main). The sequence has been trimmed to remove vector sequence and contains a minimum of 100 bases of phased value 20 or above. For more details on library preparation and sequence analysis see <http://www.genome.clemson.edu/projects/barley>. To order this clone see <http://www.genome.clemson.edu/orders> Also see Close TJ, Wing R, Kleinhoofs A, Wise R (2001) Genetically and physically anchored EST resources for barley genomics. Barley Genetics Newsletter 31:29-30. (<http://wheat.pw.usda.gov/ggpages/bgn/31/cover.html>)" 221 c 324 q 131 t

VERSION	KEYWORDS	SOURCE	ORGANISM	REFERENCE	AUTHORS	TITLE	JOURNAL	COMMENT	FEATURES	source
BF293584.1	GI:11224648	EST	Triticum turgidum	Unpublished (2000)	Contact: Olin Anderson					
BF293584.1	GI:11224648	EST	Triticum turgidum	US Department of Agriculture, Agriculture Research Service, Pacific West Area, Western Regional Research Center	800 Buchanan Street, Albany, CA 94710, USA					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	Tel: 5105959773					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	Fax: 5105959818					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	Email: oanderson@pw.usda.gov					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	Sequence have been trimmed to remove vector sequence and low quality sequence with phred score less than 20					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	Seq primer: Stratagene SK primer.					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	Location/Qualifiers					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	1. 376					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	/organism="Triticum turgidum"					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	/cultivar="Langdon-16"					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	/db_xref="taxon:4571"					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	/clone="WHE2158_E09_J18"					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	/clone_lib="Triticum turgidum L. var. durum (durum wheat)"					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	whole plant cDNA library"					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	/tissue_type="All tissues"					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	/dev_stage="Different growth stages"					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	/lab_host="E. coli SOLR"					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	/note="Vector: Lambda Uni-ZAP XR, excised phagemid;					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	Site 1: EcoRI; Site 2: XhoI; Plants were grown in a growth chamber at North Dakota State University (Kianian, Otto,					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	seedling leaf, stem, root and seed; leaf from plant at					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	fourth leaf stage; spike from pre-anthesis through 20					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	days after anthesis; flag leaf; leaf and stem tissue from					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	tillers. and root. Total RNA and poly(A) RNA were					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	prepared from each tissue and then pooled, a cDNA library					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	was made, and the cDNA clones were in vivo excised to give					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	pBluescript phagemids in the T3 Close lab (Akhunov, Chin,					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	Choi, Close, Fenton, Kianian, Otto, Simons, Zhang) at the					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	University of California, Riverside. Plasmid DNA					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	preparations and DNA sequencing were performed in the OD					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	Anderson lab (all other authors)."					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany, CA 94710, USA	Anderson lab (all other authors)."					
BF293584.1	GI:11224648	EST	Triticum turgidum	800 Buchanan Street, Albany						

```

Db 112 CGAAGAAGTCTCTCGTGGAGTCCGAGGCGCGTGTGCTGTCGACATGTACCTCCTC 171
QY 1203 cgataccagacagcgagcagcattggttggattggaattgacgaagacacccgttt 1262
Db 172 CTACTACTCTGTCGACGACAGGTTTGTTCACGAGGATTTCACGAGGAGAGATGTGGCCGG 231
QY 1263 gtgcagcttcgtctcccgaggcaatacga 1292
Db 232 GATGGGGTGGAGAGACGCGTGAAGTTTGA 261

RESULT 27
AL523270
LOCUS
DEFINITION
AL523270 LTI_NFL003_NBC3 893 bp mRNA linear EST 13-FEB-2001
prime mRNA sequence.
ACCESSION
AL523270
VERSION
AL523270.1 GI:12786763
KEYWORDS
EST.
SOURCE
human.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 893)
Li.W.B., Gruber,C., Jessee,J. and Polayes,D.
Full-length cDNA libraries and normalization
Unpublished (2001)
Contact: Genoscope
Genoscope - Centre National de Sequencage
BP 191 91006 Evry cedex - France
Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr.

FEATURES
source
1. .893
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="CS0DC001YH12"
/clone_lib="LTI_NFL003_NBC3"
/sex="male"
/tissue_type="neuroblastoma cells"
/lab_host="DH10B"
/notes="Organ: brain; Vector: pCMVSPORT 6; 1st strand cDNA
was primed with a NotI-oligo(dT) primer. Five prime end
enriched, double-stranded cDNA was digested with Not I and
cloned into the Not I and Eco RV sites of the pCMVSPORT 6
vector. Library was normalized. Library was constructed
by Life Technologies. Contact : peng liang life
Technologies, a division of Invitrogen 9800 Medical Center
Drive Rockville, Maryland 20850, USA Fax : (1) 301 610
8371 Email : fliang@lifetech.com URL :
http://fulllength.invitrogen.com"
BASE COUNT 168 a 282 c 285 g 153 t 5 others
ORIGIN

```

```

Query Match 2.8%; Score 37.2; DB 9; Length 893;
Best Local Similarity 54.8%; Pred. No. 22;
Matches 69; Conservative 2; Mismatches 55; Indels 0; Gaps 0;

QY 1215 cgcagcagcattggttggattgacgaagacccctgttggagcttcgt 1274
Db 258 CGGGGCTGCAACGCCGAGCKCTCAGGGCCGCCGAGGATGGGCTCGCTGCC 317
QY 1275 ctgccggcgcaatacgaatacggcccgctgttgcgaagtgctggaaccattgagaa 1334
Db 318 CAACCCGGGCACATTCGAGGAGTGCCACCGGAAGTGAAGGAGCTGTTCCTCCATTCAGAT 377
QY 1335 ggaagg 1340
Db 378 GGAGGG 383

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RESULT 28

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T43876
LOCUS
DEFINITION
7139 Lambda-PRL2 Arabidopsis thaliana cDNA clone 118M9T7, mRNA
sequence.
ACCESSION
T43876
VERSION
T43876.1 GI:948248
KEYWORDS
EST.
SOURCE
thale cress.
ORGANISM
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsids.
1 (bases 1 to 481)
Newman,T., deBruijn,F.J., Green,P., Keegstra,K., Kende,H., McIntosh
,L., Ohlrogge,J., Raikhel,N., Somerville,S., Thomashow,M., Retzel
,E. and Somerville,C.
Genes galore: a summary of methods for accessing results from
large-scale partial sequencing of anonymous Arabidopsis cDNA clones
Plant Physiol. 106, 1241-1255 (1994)
95148729
On Nov 29, 1993 this sequence version replaced gi:636464.
Contact: Thomas Newman
MSU-DOE Plant Research Laboratory
Michigan State University
MSU-DOE-PRL, Michigan State University, Plant Biology Bldg., E.
Lansing, MI
Tel: 517-353-0854
Fax: 517-353-9168
Email: 22313tcn@bm.cl.msu.edu
Seq primer: T7
Location/Qualifiers
1. .481
/organism="Arabidopsis thaliana"
/strain="var columbia"
/db_xref="taxon:3702"
/clone="118M9T7"
/clone_lib="Lambda-PRL2"
/notes="Vector: lambda Zip-Lox; Site_1: Sal; Site_2: Not;
Lambda PRL2 is a cDNA library derived from equal
quantities of 4 pools of mRNA. The mRNA sources were 1) 7
day germinated etiolated seedlings; 2) tissue culture
grown roots; 3) staged plants half with 24 hour light
cycle, half on 16 hr light, 8 hour dark- rosettes; 4)
same plants as 3 but aerial tissue (stems, flowers and
siliques). The vector is BRL's lambda Zip-Lox. The cDNA
inserts were directionally cloned with Sal-Not arms using
oligo dT primed cDNA."
BASE COUNT 122 a 96 c 113 g 133 t 17 others
ORIGIN

```

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Query Match 2.7%; Score 36.8; DB 10; Length 481;
Best Local Similarity 55.0%; Pred. No. 21;
Matches 71; Conservative 0; Mismatches 58; Indels 0; Gaps 0;

QY 128 tgaagtcgaaggaaggcgtgcgcgaaggaagcgaagcgtgtgttgaagacacaaaga 187
Db 47 TCAAGCTACACAGATCTCTGGAATCGAAGGCTTGANTCTGTGTATATGCGGAGGAGA 106
QY 188 atccggcggtgttactgcgcggcgttcaggcaaaatgcgcgattccacgtggtcg 247
Db 107 AATGGGTGTGTGTGTGAGCGGCGAGACTCCCAACGAATATCGCGGTGATTAAGTATTGGG 166
QY 248 aaagcgcg 256
Db 167 GAAAGAGAG 175

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RESULT 29

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AW714071/c
LOCUS
DEFINITION
AW714071 Neurospora crassa evening cDNA library Neurospora crassa
h5f02ne 5', mRNA sequence.
458 bp mRNA linear EST 25-APR-2000
cDNA clone h5f02ne 5', mRNA sequence.

```

REFERENCE
AUTHORS
TITLE
JOURNAL
MEDLINE
COMMENT
Genome Res. 11 (8), 1434-1440 (2001)
21376150
Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Class: BAC ends
High quality sequence stop: 560.
Location/Qualifiers
1. .617
/organism="Bradyrhizobium japonicum"
/strain="USDA110"
/db_xref="taxon:375"
/clone_lib="B. japonicum BAC library"
/lab_host="E. coli"
/note="Vector: pIndigo536; Site_1: HindIII"
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ORIGIN
Query Match 2.7%; Score 36.4; DB 12; Length 617;
Best Local Similarity 47.9%; Pred. No. 30;
Matches 103; Conservative 0; Mismatches 112; Indels 0; Gaps 0;
QY 1101 cgtcaacggcgagcgccatggtgcgattggtactacgagcggtgacccctt 1160
Db 328 CATCATGACGGCGGCGGACCATCGCGCGGTCGAGGCGCATGAGCGGGCGGT 387
QY 1161 ggtatctgcccacccctgttttgcgcgatttaactgcgcgataccgacgagcgca 1220
Db 388 CGACTACATTCGAGCCGTTCAAGCTCAGCGTCGTCATTCCTGCTCGAGCGCGGT 447
QY 1221 ggcattgggtgtcttggaattgacgaagaagacctgttgcagcttgcgttcgccc 1280
Db 448 CGCGTTCGGCGCGCTGCACCTGCGAGATGNGAGCTGGCGGTCCGCGGCGGAC 507
QY 1281 gggcaatacgaatacggcgctgtgttgccaaag 1315
Db 508 CCTCGAGCTCGAAGCCGCCACCGGGAGCTCGAGG 542
RESULT 31
CNS02UAR/c
LOCUS
DEFINITION
CNS02UAR 967 bp DNA linear GSS 15-MAY-2000
Tetraodon nigroviridis genome survey sequence PUC-ori end of clone
166L18 of library G from Tetraodon nigroviridis, genomic survey
sequence.
AL214236
AL214236.1 GI:7873055
GSS; genome survey sequence.
Tetraodon nigroviridis
Tetraodon nigroviridis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
Tetraodontidae; Tetraodon.
1 (bases 1 to 967)
Roest-Crollius.H., Jaillon.O., Dasilva.C., Fizames.C., Fisher.C.,
Bouneau.L., Billault.A., Quetier.F., Saurin.W., Bernot.A. and
Weissenbach.J.
Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
Unpublished
2 (bases 1 to 967)
Roest-Crollius.H., Jaillon.O., Dasilva.C., Bouneau.L., Fisher.C.,

REFERENCE
AUTHORS
TITLE
JOURNAL
MEDLINE
COMMENT
Genome Res. 11 (8), 1434-1440 (2001)
21376150
Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Class: BAC ends
High quality sequence stop: 560.
Location/Qualifiers
1. .617
/organism="Bradyrhizobium japonicum"
/strain="USDA110"
/db_xref="taxon:375"
/clone_lib="B. japonicum BAC library"
/lab_host="E. coli"
/note="Vector: pIndigo536; Site_1: HindIII"
BASE COUNT 108 a 193 c 216 g 98 t 2 others
ORIGIN
Query Match 2.7%; Score 36.4; DB 12; Length 617;
Best Local Similarity 47.9%; Pred. No. 30;
Matches 103; Conservative 0; Mismatches 112; Indels 0; Gaps 0;
QY 1101 cgtcaacggcgagcgccatggtgcgattggtactacgagcggtgacccctt 1160
Db 328 CATCATGACGGCGGCGGACCATCGCGCGGTCGAGGCGCATGAGCGGGCGGT 387
QY 1161 ggtatctgcccacccctgttttgcgcgatttaactgcgcgataccgacgagcgca 1220
Db 388 CGACTACATTCGAGCCGTTCAAGCTCAGCGTCGTCATTCCTGCTCGAGCGCGGT 447
QY 1221 ggcattgggtgtcttggaattgacgaagaagacctgttgcagcttgcgttcgccc 1280
Db 448 CGCGTTCGGCGCGCTGCACCTGCGAGATGNGAGCTGGCGGTCCGCGGCGGAC 507
QY 1281 gggcaatacgaatacggcgctgtgttgccaaag 1315
Db 508 CCTCGAGCTCGAAGCCGCCACCGGGAGCTCGAGG 542
RESULT 31
CNS02UAR/c
LOCUS
DEFINITION
CNS02UAR 967 bp DNA linear GSS 15-MAY-2000
Tetraodon nigroviridis genome survey sequence PUC-ori end of clone
166L18 of library G from Tetraodon nigroviridis, genomic survey
sequence.
AL214236
AL214236.1 GI:7873055
GSS; genome survey sequence.
Tetraodon nigroviridis
Tetraodon nigroviridis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
Tetraodontidae; Tetraodon.
1 (bases 1 to 967)
Roest-Crollius.H., Jaillon.O., Dasilva.C., Fizames.C., Fisher.C.,
Bouneau.L., Billault.A., Quetier.F., Saurin.W., Bernot.A. and
Weissenbach.J.
Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
Unpublished
2 (bases 1 to 967)
Roest-Crollius.H., Jaillon.O., Dasilva.C., Bouneau.L., Fisher.C.,

REFERENCE
AUTHORS
TITLE
JOURNAL
MEDLINE
COMMENT
Genome Res. 11 (8), 1434-1440 (2001)
21376150
Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Class: BAC ends
High quality sequence stop: 560.
Location/Qualifiers
1. .617
/organism="Bradyrhizobium japonicum"
/strain="USDA110"
/db_xref="taxon:375"
/clone_lib="B. japonicum BAC library"
/lab_host="E. coli"
/note="Vector: pIndigo536; Site_1: HindIII"
BASE COUNT 108 a 193 c 216 g 98 t 2 others
ORIGIN
Query Match 2.7%; Score 36.4; DB 12; Length 617;
Best Local Similarity 47.9%; Pred. No. 30;
Matches 103; Conservative 0; Mismatches 112; Indels 0; Gaps 0;
QY 1101 cgtcaacggcgagcgccatggtgcgattggtactacgagcggtgacccctt 1160
Db 328 CATCATGACGGCGGCGGACCATCGCGCGGTCGAGGCGCATGAGCGGGCGGT 387
QY 1161 ggtatctgcccacccctgttttgcgcgatttaactgcgcgataccgacgagcgca 1220
Db 388 CGACTACATTCGAGCCGTTCAAGCTCAGCGTCGTCATTCCTGCTCGAGCGCGGT 447
QY 1221 ggcattgggtgtcttggaattgacgaagaagacctgttgcagcttgcgttcgccc 1280
Db 448 CGCGTTCGGCGCGCTGCACCTGCGAGATGNGAGCTGGCGGTCCGCGGCGGAC 507
QY 1281 gggcaatacgaatacggcgctgtgttgccaaag 1315
Db 508 CCTCGAGCTCGAAGCCGCCACCGGGAGCTCGAGG 542
RESULT 31
CNS02UAR/c
LOCUS
DEFINITION
CNS02UAR 967 bp DNA linear GSS 15-MAY-2000
Tetraodon nigroviridis genome survey sequence PUC-ori end of clone
166L18 of library G from Tetraodon nigroviridis, genomic survey
sequence.
AL214236
AL214236.1 GI:7873055
GSS; genome survey sequence.
Tetraodon nigroviridis
Tetraodon nigroviridis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
Tetraodontidae; Tetraodon.
1 (bases 1 to 967)
Roest-Crollius.H., Jaillon.O., Dasilva.C., Fizames.C., Fisher.C.,
Bouneau.L., Billault.A., Quetier.F., Saurin.W., Bernot.A. and
Weissenbach.J.
Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
Unpublished
2 (bases 1 to 967)
Roest-Crollius.H., Jaillon.O., Dasilva.C., Bouneau.L., Fisher.C.,

EcoRI; See: Bell-Perdersen, D., et al. PNAS 93:13096, 1996. 5' end of cDNA cloned into XbaI site of pBluescript; 3' end of cDNA cloned into EcoRI site of pBluescript"

BASE COUNT	97 a	150 c	76 g	84 t
ORIGIN				
Query Match		2.7%	Score 36.2;	DB 9; Length 407;
Best Local Similarity		48.3%;	Prsd. NO. 28;	
Matches 101; Conservative		0;	Mismatches 108;	Indels 0; Gaps 0;

[illegible]

RESULT	34
BF428970	
LOCUS	
DEFINITION	
BF428970	407 bp mRNA linear EST 29-NOV-2000
WHEI1712_E10_J20ZS wheat heat stressed spike cDNA library Trilicum aestivum cDNA clone WHEI1712 E10 J20.	mRNA sequence.

ACCESSION	BF428970	
VERSION	BF428970.1	GI:11440887
KEYWORDS	EST.	
SOURCE	bread wheat.	
ORGANISM	Triticum aestivum	
REFERENCE	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae	
AUTHORS	1 (bases 1 to 407)	
TITLE	Triticaceae; Triticum.	
JOURNAL	Anderson, O.D., Chao, S., Choi, D.W., Close, T.J., Fenton, R.D., Han, P.S., Hsia, C.C., Kang, Y., Lazo, G.R., Malatrasi, M., Miller, R., Nguyen, H.T., Rausch, C.J., Seaton, C.L., Tong, J.C. and Zhang, D.	
	The structure and function of the expressed portion of the wheat genomes - Heat stressed spike cDNA library	
	Unpublished (2000)	

COMMENT

Contact: Olin Anderson
US Department of Agriculture, Agriculture Research Service, Pacific
West Area, Western Regional Research Center
800 Buchanan Street, Albany, CA 94710, USA
Tel: 5105959773
Fax: 5105959818
Email: oanderson@pw.usda.gov

Sequence have been trimmed to remove vector sequence and low
quality sequence with phred score less than 20

Seq primer: StrataGene SK primer.

FEATURES	Location/Qualifiers
1. .407	/organism="Triticum aestivum"
	/cultivar="Chinese Spring"
	/db_xref="taxon:4565"
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	/clone_lib="wheat heat stressed spike cDNA library"
	/tissue_type="Whole spike"
	/dev_stage="Spikes at 5, 10, 15 and 20 days after anthesis"
	/lab_host="E. coli SOLR"
	/note="Vector: lambda Uni-ZAP XR, excised phagemid; Site_1: EcoRI; Site_2: XhoI; Spikes at 5, 10, 15 and 20 days after anthesis were heat stressed under two conditions at Texas Tech University (D. Zhang in HT Nguyen

lab): (1) at 38 C for 4 hours and (2) 5 days of cyclic treatment at 38 C for 4 hours. Total RNA and poly(A)⁺ RNA were prepared, a cDNA library was made, and the cDNA clones were *in vivo* excised to give Bluescript phagemids at the University of California, Riverside, Malatrasí at the University of California, Riverside, plasmid DNA preparations and DNA sequencing were performed in the Anderson lab (all other authors).

[illegible]

RESULT 35

	BG262553	linear	mRNA	EST 16-FEB-2001
LOCUS	BG262553	460 bp	wheat spike cDNA library	Triticum
DEFINITION	WHE90937.DL0_IG19ZS	Wheat 5-15 DAP	cDNA clone whe90937 n10 G19.	mRNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE

ORGANISM

Triticum aestivum
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
Triticeae; Triticum

REFERENCE

AUTHORS Anderson, O.D., Chao, S., Choi, D.W., Close, T.J., Fenton, R.D., Han, P.S., Hsia, C.C., Kang, Y., Lazo, G.R., Miller, R., Rausch, C.J., Seaton, C.L. and Tong, J.C.

TITLE The structure and function of the wheat genomes - 5-15 DAP spike cDNA library unpublished (2000)

JOURNAL

COMMENT

US Department of Agriculture, Agriculture Research Service, Facilities
West Area, Western Regional Research Center
800 Buchanan Street, Albany, CA 94710, USA
Tel: 5105959773
Fax: 5105505818

Email: oandern@pw.usda.gov
 Sequence have been trimmed to remove vector sequence and low
 quality sequence with phred score less than 20
 Sac primer: StrataGene SK primer.

FEATURES SOURCES

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1. .460
/organism="Triticum aestivum"
/cultivar="Chinese Spring"
/db_xref="taxon:4565"
/clone="WHE0937_D10_G19"
/clone_lib="Wheat_5-15_DAP_spike_cdna_library"
/tissue_type="Spike"
/dev_stage="Adult plant"
/lab_host="E. coli SOLR"
/note="vector: Lambda Uni-ZAP XR, excised phagemid;
site_1: EcoRI; site_2: XhoI; plants were grown in the
greenhouse. Spikes at 5, 10 and 15 DAP were harvested,
total RNA and poly(A) RNA were prepared, a cDNA library

```

was made, and the cDNA clones were in vivo excised to give pBluescript phagemids in the TJ Close lab (Choi, Close, Fenton) at the University of California, Riverside. Plasmid DNA preparations and DNA sequencing were performed in the OD Anderson lab (all other authors).

```

BASE COUNT      131 a  117 c  118 g  94 t
ORIGIN

Query Match      2.7%; Score 36.2; DB 10; Length 460;
Best Local Similarity 50.3%; Pred. No. 30;
Matches 89; Conservative 0; Mismatches 88; Indels 0; Gaps 0;

QY 152 tcaaaagggccaaagtctgtttgaagacaaaagaatccggcggtgtttactgcgc 211
    |||||
DB 191 TCAAGCATGGTGACAAAGATCTGGACGACCAACAAAGGCTCTCTTTGGGACACACAGTGC 250
    |||||
QY 212 cggttcaggcaaaatcgccgattccacgtgcccgaagcgcgctacttcagtcagtcg 271
    |||||
DB 251 AACTGGCAATCAAAATCTCTGGCATTCACCTGGTGTGACAAAGGTTCCAAAGCCATGCAGCGC 310
    |||||
QY 272 tgattgcgttggaagcaacgcagcaaatcgagtttgaaacgtacgcacctgaagcgc 328
    |||||
DB 311 AGGTACACAGCTGGGTACTACAACTTACATGATAGACGGCTACAGACCCATAGCAC 367
    |||||

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RESULT 36
BE590713
LOCUS      BE590713      460 bp      mRNA      linear      EST 18-AUG-2000
DEFINITION WHE0856_A07_A142S Wheat 20-45 DAP spike cDNA library. Triticum aestivum cDNA clone WHE0856_A07_A14, mRNA sequence.
ACCESSION BE590713
VERSION    BE590713.1 GI:9845786
KEYWORDS   EST.
SOURCE     bread wheat.
ORGANISM   Triticum aestivum
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
            1 (bases 1 to 460)
            P.S., Hsiao, C.C., Kang, Y., Lazo, G.R., Miller, R., Rausch, C.J.,
            Seaton, C.L. and Tong, J.C.
            The structure and function of the expressed portion of the wheat
            genomes - 20-45 DAP spike cDNA library
            Unpublished (2000)
            Contact: Olin Anderson
            US Department of Agriculture, Agriculture Research Service, Pacific
            West Area, Western Regional Research Center
            800 Buchanan Street, Albany, CA 94710, USA
            Tel: 5105955773
            Fax: 5105959518
            Email: andersn@pw.usda.gov
            Sequence have been trimmed to remove vector sequence and low
            quality sequence with phred score less than 20
            Seq primer: Stratagene SK primer.
            Location/Qualifiers
                1..460
                /organism="Triticum aestivum"
                /cultivar="Chinese Spring"
                /db_xref="taxon:4565"
                /clone="WHE0856_A07_A14"
                /tissue_type="Spike and seed"
                /dev_stage="Adult plant"
                /lab_host="E. coli SOLR"
                /note="Vector: Lambda Uni-ZAP XR, excised phagemid;
                Site 1: EcoRI; Site 2: XhoI; Plants were grown in the
                greenhouse. Spikes at 20 DAP and seeds at 30 to 45 DAP
                were harvested. total RNA and poly(A) RNA were prepared, a
                cDNA library was made, and the cDNA clones were in vivo
                excised to give pBluescript phagemids in the TJ Close lab

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FEATURES
source

(Choi, Close, Fenton) at the University of California, Riverside. Plasmid DNA preparations and DNA sequencing were performed in the OD Anderson lab (all other authors).

```

BASE COUNT      129 a  117 c  114 g  100 t
ORIGIN

Query Match      2.7%; Score 36.2; DB 10; Length 460;
Best Local Similarity 50.3%; Pred. No. 30;
Matches 89; Conservative 0; Mismatches 88; Indels 0; Gaps 0;

QY 152 tcaaaagggccaaagtctgtttgaagacaaaagaatccggcggtgtttactgcgc 211
    |||||
DB 227 TCAAGCATGGTGACAAAGATCTGGACGACCAACAAAGGCTCTCTTTGGGACACACAGTGC 286
    |||||
QY 212 cggttcaggcaaaatcgccgattccacgtgcccgaagcgcgctacttcagtcagtcg 271
    |||||
DB 287 AACTGGCAATCAAAATCTCTGGCATTCACCTGGTGTGACAAAGGTTCCAAAGCCATGCAGCGC 346
    |||||
QY 272 tgattgcgttggaagcaacgcagcaaatcgagtttgaaacgtacgcacctgaagcgc 328
    |||||
DB 347 AGGTACACAGCTGGGTACTACAACTTACATGATAGACGGCTACAGACCCATAGCAC 403
    |||||

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RESULT 37
AA614629/c
LOCUS      AA614629      505 bp      mRNA      linear      EST 16-OCT-1997
DEFINITION np54b11.s1 NCI-CGAP.Br1.1 Homo sapiens cDNA clone IMAGE:1130109 3'
            similar to gb:J04456 GALECTIN-1 (HUMAN);, mRNA sequence.
ACCESSION AA614629
VERSION    AA614629.1 GI:2466825
KEYWORDS   EST.
SOURCE     human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
            1 (bases 1 to 505)
            NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
            National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
            Unpublished (1997)
            Contact: Robert Strausberg, Ph.D.
            Email: cgapbs-r@mail.nih.gov
            Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
            Emmert-Buck, M.D., Ph.D.
            cDNA Library Preparation: M. Bento Soares, Ph.D.
            DNA Sequencing by: Greg Lennon, Ph.D.
            Cloning Strategy: Washington University Genome Sequencing Center
            Clone distribution: NCI-CGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LNL at:
            www.bio.lnlnl.gov/bbrp/image/image.html
            Insert Length: 549 Std Error: 0.00
            Seq primer: -40ml3 fwd. ET from Amersham
            High quality sequence stop: 388.
            Location/Qualifiers
                1..505
                /organism="Homo sapiens"
                /db_xref="taxon:9606"
                /clone="IMAGE:1130109"
                /clone_lib="NCI-CGAP.Br1.1"
                /sex="female, pooled"
                /tissue_type="breast"
                /lab_host="DH10B"
                /note="Vector: pT7T3D-Pac (Pharmacia) with a modified
                polylinker; 1st strand cDNA was prepared from pooled bulk
                breast tumor tissue, and was then primed with a Not I -
                oligo(dT) primer. Double-stranded cDNA was ligated to Eco
                RI adaptors (Pharmacia), digested with Not I and cloned
                into the Not I and Eco RI sites of the modified pT7T3
                vector. Library is not normalized. (The normalized
                version of this library is NCI-CGAP.Br2.) Library was
                constructed by Bento Soares and M. Fatima Bonaldo."

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FEATURES
source

100

1


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Query Match      2.7%; Score 36.2; DB 10; Length 625;
Best Local Similarity 50.3%; Pred. No. 35;
Matches 89; Conservative 0; Mismatches 88; Indels 0; Gaps 0;

QY 152 tcaaaagggcccaagtgtgtttgaagacaaagaaatccggcggtgtgtttactgcgc 211
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 252 TCAGCATGGTGCACAGATCTTGGACGAAGCAACAAAGGCTTCTTGGGACACACAGTCG 311
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 212 cggcttcagggaacatcccgatccacgtggcgaaagcgcggtacttcagtcagtcg 271
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 312 AACTGCAATCAAAATCTTGGCATCTTGGCATCTTGGCATCTTGGCATCTTGGCATCTTGGCAT 371
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 272 tgattgcggtgaaggaacacacacacacacacacacacacacacacacacacacacac 328
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 372 AGGTACAGCTGGGTACTACAACTTACATGATAGAGACGGCTACAGACCCCATAGCAC 428
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

RESULT 40
BG605187
LOCUS      BG605187      639 bp      mRNA      linear      EST 16-APR-2001
DEFINITION Wheat pre-anthesis spike cDNA library Triticum
aestivum CDNA clone WHE2328_F06_L12, mRNA sequence.
ACCESSION      BG605187
VERSION      BG605187.1
KEYWORDS      EST.
SOURCE      bread wheat.
ORGANISM      Triticum aestivum
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
; Triticaceae; Triticum.
1 (bases 1 to 639)
Anderson, O.D., Chao, S., Choi, D.W., Close, T.J., Fenton, R.D., Han
, P.S., Hsia, C.C., Kang, Y., Lazo, G.R., Miller, R., Rausch, C.J.,
Seaton, C.L. and Tong, J.C.
The structure and function of the expressed portion of the wheat
genomes - Pre-anthesis spike cDNA library
Unpublished (2000)
Contact: Olin Anderson
US Department of Agriculture, Agriculture Research Service, Pacific
West Area, Western Regional Research Center
800 Buchanan Street, Albany, CA 94710, USA
Tel: 5105959773
Fax: 5105959818
Email: oanderson@pw.usda.gov
Sequence have been trimmed to remove vector sequence and low
quality sequence with phred score less than 20
Seq primer: Stratagene SK primer.
FEATURES
source
1..639
Location/Qualifiers
/organism="Triticum aestivum"
/cultivar="Chinese Spring"
/db_xref="taxon:4565"
/clone="WHE2328_F06_L12"
/clone_lib="Wheat pre-anthesis spike cDNA library"
/tissue_type="Spike before anthesis"
/dev_stage="Adult plant"
/lab_host="E. coli SOLR"
/note="Vector: Lambda Uni-ZAP XR, excised phagemid;
Site 1: EORI; Site 2: XhoI; Plants were grown in the
greenhouse. Whole spike with awns trimmed, white, green
and yellow anther were collected and total RNA, and
poly(A) RNA were prepared, a cDNA library was made, and
the cDNA clones were in vivo excised to give pBluescript
phagemids in the TJ Close lab (Choi, Close, Penton) at
the University of California, Riverside. Plasmid DNA
preparations and DNA sequencing were performed in the OD
Anderson lab (all authors)."
BASE COUNT      182 a 165 c 160 g 132 t
ORIGIN

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Query Match      2.7%; Score 36.2; DB 10; Length 639;
Best Local Similarity 50.3%; Pred. No. 35;

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Matches 89; Conservative 0; Mismatches 88; Indels 0; Gaps 0;

QY 152 tcaaaagggcccaagtgtgtttgaagacaaagaaatccggcggtgtgtttactgcgc 211
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 196 TCAGCATGGTGCACAGATCTTGGACGAAGCAACAAAGGCTTCTTGGGACACACAGTCG 255
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 212 cggcttcagggaacatcccgatccacgtggcgaaagcgcggtacttcagtcagtcg 271
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 256 AACTGCAATCAAAATCTTGGCATCTTGGCATCTTGGCATCTTGGCATCTTGGCATCTTGGCAT 315
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 272 tgattgcggtgaaggaacacacacacacacacacacacacacacacacacacacacac 328
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 316 AGGTACAGCTGGGTACTACAACTTACATGATAGAGACGGCTACAGACCCCATAGCAC 372
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

RESULT 41
AL556080
LOCUS      AL556080      943 bp      mRNA      linear      EST 16-FEB-2001
DEFINITION LTI_NFL006_PL2 Homo sapiens CDNA clone CSODK010YJ05 5
prime, mRNA sequence.
ACCESSION      AL556080
VERSION      AL556080.1
KEYWORDS      EST.
SOURCE      human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 943)
Li, W.B., Gruber, C., Jessee, J. and Pollayes, D.
Full-length cDNA libraries and normalization
Unpublished (2001)
Contact: Genoscope
Genoscope - Centre National de Sequencage
BP 191 91006 EVRY cedex - France
Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr.
FEATURES
source
1..943
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="CSODK010YJ05"
/clone_lib="LTI_NFL006_PL2"
/tissue_type="placenta"
/note="Vector: pCMVSPORT 6; Site 1: NotI; 1st strand cDNA
was primed with a NotI-oligo(dT) primer. Five prime end
enriched, double-stranded cDNA was digested with Not I and
cloned into the Not I and Eco RV sites of the pCMVSPORT 6
vector. Library was normalized. Library was constructed by
Life Technologies, Contact : Feng Liang Life Technologies,
a division of Invitrogen 9800 Medical Center Drive
Rockville, Maryland 20850, USA Fax : (1) 301 610 8371
Email : fliang@lifetech.com URL :
http://fulllength.invitrogen.com"
BASE COUNT      185 a 293 c 297 g 152 t 16 others
ORIGIN

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Query Match      2.7%; Score 36.2; DB 9; Length 943;
Best Local Similarity 59.6%; Pred. No. 42;
Matches 59; Conservative 1; Mismatches 39; Indels 0; Gaps 0;

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```

QY 1242 ggacgaagaagacatcgcttctgtgcagcttcgtcccgaggcaataacacgacgc 1301
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 345 GCCGCCCGAGGATGGGGCTCGGCTGCCCAACCCGGGCACATCCGAGGATGCCA 404
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 1302 gctgtgtcgaaatgctggaaacattgagaagaagg 1340
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 405 CCGGAGTGCAGAGAGCTGTTCCTCCATCAATGARGG 443
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

RESULT 42
AV939450
LOCUS      AV939450      489 bp      mRNA      linear      EST 18-JAN-2002
DEFINITION AV939450 K. Sato unpublished cDNA library, strain H602 adult,

```

AQ161443
 AQ161443.1 GI:3557844
 GSS.
 Magnaporthe grisea.
 Magnaporthe grisea
 Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes;
 Sordariomycetes incertae sedis; Magnaporthaceae; Magnaporthe.
 1 (bases 1 to 612)
 Yu, Y., Zhu, H., Boyd, C.A., Gaudette, B., Gayle, A., Kingsbury, R.,
 Phillips, K., Sasinowski, M., Wang, R.A. and Dean, R.A.
 A BAC End Sequencing Framework to Sequence the *Magnaporthe grisea*
 Genome

REFERENCES

1 (bases 1 to 779)

1 (bases 1 to 779)

1. The first part of the paper is devoted to a general discussion of the problem of the existence of solutions of the system of equations